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International Researcher IDs

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Publons / Web Of Science ResearcherID: A-5099-2019

ScopusID: 60030785

Yoksis Researcher ID: 36748

Education Information

Post Doctorate of Medicine, Erciyes University, Tıp Fakültesi, Çocuk Hematoloji Ve Onkoloji Bilim Dalı, Turkey 2007 - 2012

Expertise In Medicine, Selcuk University, School Of Medicine, Çocuk Sağlığı Ve Hastalıkları, Turkey 2001 - 2007

Undergraduate, Selcuk University, School Of Medicine, Turkey 1995 - 2001

Foreign Languages

English, C1 Advanced

Certificates, Courses and Trainings

Health&Medicine, Deney Hayvanları Kullanım sertifikası, Erciyes Üniversitesi, 2012

Health&Medicine, Hemofili Akademisi, Edinburg, İngiltere, 2011

Education Management and Planning, Tıp Fakültesi Öğrencilerine yönelik Eğitim Sertifikası, Erciyes Üniversitesi Tıp Fakültesi, 2011

Health&Medicine, Lösemide MRD, Charite Univeritaetsmedizin Berlin Almanya, 2010

Health&Medicine, İyi Klinik Uygulamalar, TC Sağlık Bakanlığı/ Türk Hematoloji Derneği, 2010

Health&Medicine, ÇİLYAD (Çocuk İleri Yaşam Desteği) modülü Eğitici Sertifikası, TC Sağlık Bakanlığı/ Konya İl Sağlık müdürlüğü, 2006

Dissertations

Expertise In Medicine, ÇOCUKLUK ÇAĞI AKUT LENFOBLASTİK LÖSEMİ HASTALARINDA ÇÖZÜNEBİLİR CD 40 VE CD 40 LİGAND SEVİYESİNİN ÖNEMİ, Erciyes Üniversitesi, Tıp Fakültesi, Çocuk Hematoloji Ve Onkoloji Bilim Dalı, 2012

Expertise In Medicine, ÇOCUKLUK ÇAĞI AKUT LÖSEMİ VE HODGKİN-DIŞI LENFOMA HASTALARINDA SİSTATİN-C' NİN

Research Areas

Health Sciences, Medicine, Internal Medicine Sciences, Child Health and Diseases, Pediatrics Infectious Diseases, Internal Diseases , Hematology, Basic Sciences, Life Sciences, Molecular Biology and Genetics, Genetic Disorders

Academic Titles / Tasks

Professor, Erciyes University, Tıp Fakültesi, Dahili Tıp Bil., 2021 - Continues

Associate Professor, Erciyes University, Tıp Fakültesi, Dahili Tıp Bil., 2015 - 2021

Assistant Professor, Erciyes University, Tıp Fakültesi, Dahili Tıp Bil., 2011 - 2015

Research Assistant, Erciyes University, Tıp Fakültesi, Dahili Tıp Bil., 2007 - 2011

Research Assistant, Selcuk University, School Of Medicine, Çocuk Sağlığı Ve Hastalıkları Anabilim Dalı, 2001 - 2007

Courses

ÇOCUKLARDA PARAZİTER HASTALIKLAR, Postgraduate, 2012 - 2013

Published journal articles indexed by SCI, SSCI, and AHCI

- I. **Efficacy of T-cell assays for the diagnosis of primary defects in cytotoxic lymphocyte exocytosis**
Chiang S. C. C., Covill L. E., Tesi B., Campbell T. M., Schlums H., Nejati-Zendegani J., Mördrup K., Wood S., Theorell J., Sekine T., et al.
Blood, vol.144, no.8, pp.873-887, 2024 (SCI-Expanded)
- II. **Pyruvate kinase deficiency in 29 Turkish patients with two novel intronic variants**
GÖK V., LEBLEBİSATAN G., Gokcebay D. G., GÜLER S., Dogan M. E., Bozdogan S. T., Yozgat A. K., ÖZCAN A., Sahinoglu E. P., Tokgoz H., et al.
BRITISH JOURNAL OF HAEMATOLOGY, vol.205, no.1, pp.236-242, 2024 (SCI-Expanded)
- III. **Identification of the molecular etiology in rare congenital hemolytic anemias using next-generation sequencing with exome-based copy number variant analysis**
IŞIK E., AYDINOK Y., Albayrak C., Durmus B., Karakas Z., Orhan M. F., SARPER N., Aydin S., ÜNAL S., Oymak Y., et al.
EUROPEAN JOURNAL OF HAEMATOLOGY, vol.113, no.1, pp.82-89, 2024 (SCI-Expanded)
- IV. **Alantolactone ameliorates graft versus host disease in mice.**
Odabas G. P., Aslan K., Suna P. A., Kendirli P. K., Erdem Ş., Çakır M., Özcan A., Yılmaz E., Karakukcu M., Donmez-Altuntas H., et al.
International immunopharmacology, vol.128, pp.111560, 2024 (SCI-Expanded)
- V. **A case of autoimmune lymphoproliferative syndrome with a novel de novo FAS variant**
ÖZÇELİK F., ASLAN K., Gok V., Ari M. B., ÖZCAN A., EKEN A., Ünal E., ÖZKUL Y., DÜNDAR M.
Pediatric Hematology and Oncology, vol.41, no.4, pp.301-309, 2024 (SCI-Expanded)
- VI. **A Novel Biallelic LCK Variant Resulting in Profound T-Cell Immune Deficiency and Review of the Literature**
Lanz A., Erdem S., ÖZCAN A., CEYLANER G., CANSEVER M., CEYLANER S., Conca R., Magg T., Acuto O., Latour S., et al.
Journal of Clinical Immunology, vol.44, no.1, 2024 (SCI-Expanded)
- VII. **JAK inhibitor treatment for inborn errors of JAK/STAT signaling: An ESID/EBMT-IEWP retrospective study**
Fischer M., Olbrich P., Hadjadj J., Aumann V., Bakhtiar S., Barlogis V., von Bismarck P., Bloomfield M., Booth C.,

Buddingh E. P., et al.

Journal of Allergy and Clinical Immunology, vol.153, no.1, pp.275, 2024 (SCI-Expanded)

- VIII. **C-terminal variants in *CDC42* drive type I interferon-dependent autoinflammation in NOCARH syndrome reversible by ruxolitinib**
Kapp F. G., Kretschmer S., Beckmann C. C., Wäsch L., Molitor A., Carapito R., Schubert M., Lucas N., Conrad S., Poignant S., et al.
CLINICAL IMMUNOLOGY, vol.256, 2023 (SCI-Expanded)
- IX. **Defective Treg generation and increased type 3 immune response in leukocyte adhesion deficiency 1**
Erdem S., HASKOLOĞLU Z. Ş., Haliloglu Y., Celikzencir H., Arik E., Keskin O., Eltan S. B., YÜCEL E., CANATAN H., AVCILAR H., et al.
Clinical Immunology, vol.253, 2023 (SCI-Expanded)
- X. **Monogenic early-onset lymphoproliferation and autoimmunity: Natural history of STAT3 gain-of-function syndrome.**
Leiding J. W., Vogel T. P., Santarlas V. G. J., Mhaskar R., Smith M. R., Carisey A., Vargas-Hernandez A., Silva-Carmona M., Heeg M., Rensing-Ehl A., et al.
The Journal of allergy and clinical immunology, vol.151, no.4, pp.1081-1095, 2023 (SCI-Expanded)
- XI. **T-cell/histiocyte-rich large B-cell lymphoma in a patient with a novel frameshift MSH6 mutation**
ÇEKİÇ Ş., AYDIN F., Karali Y., SEVİNİR B. B., CANÖZ Ö., Boztug K., ÜNAL E., KILIÇ GÜLTEKİN S. Ş.
PEDIATRIC BLOOD & CANCER, vol.70, no.3, 2023 (SCI-Expanded)
- XII. **Hypomorphic RAG deficiency: impact of disease burden on survival and thymic recovery argues for early diagnosis and HSCT**
Schuetz C., Gerke J., Ege M., Walter J., Kusters M., Worth A., Kanakry J., Dimitrova D., Wolska-Kuśnierz B., Chen K., et al.
Blood, vol.141, no.7, pp.713-724, 2023 (SCI-Expanded)
- XIII. **Human genetic defects in SRP19 and SRPRA cause severe congenital neutropenia with distinctive proteome changes.**
Linder M. I., Mizoguchi Y., Hesse S., Csaba G., Tatematsu M., Łyszkiwicz M., Zietara N., Jeske T., Hastreiter M., Rohlf M., et al.
Blood, vol.141, no.6, pp.645-658, 2023 (SCI-Expanded)
- XIV. **Human CARMIL2 deficiency underlies a broader immunological and clinical phenotype than CD28 deficiency**
Levy R., Gothe F., Momenilandi M., Magg T., Materna M., Peters P., Raedler J., Philippot Q., Rack-Hoch A. L., Langlais D., et al.
JOURNAL OF EXPERIMENTAL MEDICINE, vol.220, no.2, 2023 (SCI-Expanded)
- XV. **Antibody Response against Vaccine Antigens in Children after TCR $\alpha\beta$ -Depleted Haploidentical Stem Cell Transplantation: Is It Similar to That in Recipients with Fully Matched Donors?**
Kondolot M., Yilmaz E., Sahin N. E., Ozcan A., Kaynar L., Unal E., Karakukcu M.
Transplantation and cellular therapy, vol.29, no.2, 2023 (SCI-Expanded)
- XVI. **Haploidentical Antibody Response against Vaccine Antigens in Children after TCR $\alpha\beta$ -Depleted Haploidentical Stem Cell Transplantation: Is It Similar to That in Recipients with Fully Matched Donors?**
KONDOLOT M., YILMAZ E., Sahin N. E., ÖZCAN A., KAYNAR L., ÜNAL E., KARAKÜKCÜ M.
TRANSPLANTATION AND CELLULAR THERAPY, no.2, 2023 (SCI-Expanded)
- XVII. **Aceruloplasminemia presenting with microcytic anemia in a Turkish boy due to a novel pathogenic variant**
Gok V., Ozcan A., Ozer S., Karaman F., Aykutlu E., Yilmaz E., Karakukcu M., Bisgin A., Unal E.
PEDIATRIC HEMATOLOGY AND ONCOLOGY, vol.40, no.7, pp.673-681, 2023 (SCI-Expanded)
- XVIII. **Treatment of Infantile Fibrosarcoma in the Era of Targeted Therapies**
Yildirim U. M., Kebudi R., Asarcikli F., Sozmen B. O. F. L. A. Z., ÜNAL E., ÖZCAN A., Zulfikar B.
UHOD-ULUSLARARASI HEMATOLOJİ-ONKOLOJİ DERGİSİ, no.4, pp.240-247, 2023 (SCI-Expanded)
- XIX. **Assessment of extracorporeal photopheresis related cell damage.**

Samur B. M., KARAKÜKCÜ Ç., ÖZCAN A., ÜNAL E., YILMAZ E., KARAKÜKCÜ M.

Transfusion and apheresis science : official journal of the World Apheresis Association : official journal of the European Society for Haemapheresis, vol.61, no.6, pp.103472, 2022 (SCI-Expanded)

- XX. **A pilot study for treatment of severe COVID-19 pneumonia by aerosolized formulation of convalescent human immune plasma exosomes (ChipEXO (TM))**
GÜL F., GÖNEN Z. B., Jones O. Y., Taşlı N. P., ZARARSIZ G., ÜNAL E., ÖZDARENDELİ A., Şahin F., EKEN A., YILMAZ S., et al.
FRONTIERS IN IMMUNOLOGY, vol.13, 2022 (SCI-Expanded)
- XXI. **Central Nervous System Fungal Infections in Children With Leukemia and Undergoing Hematopoietic Stem Cell Transplantation: A Retrospective Multicenter Study**
Karaman S., Kebudi R., Kizilocak H., Karakas Z., Demirag B., SEZGİN EVİM M., Yaralı N., KAYA Z., Karagun B. S., Aydogdu S., et al.
JOURNAL OF PEDIATRIC HEMATOLOGY ONCOLOGY, vol.44, no.8, 2022 (SCI-Expanded)
- XXII. **Evaluation of primary care physicians' approaches to hemophilia and bleeding disorders: a questionnaire survey**
Samur B. M., Samur T. G., Çiflikli F., Özcan A., Gök V., Soykan R., Soytürk F., Kılıç Ö., Kandur M., Kandemir R., et al.
BLOOD COAGULATION & FIBRINOLYSIS, vol.33, no.7, pp.381-388, 2022 (SCI-Expanded)
- XXIII. **Characterization of cord blood CD3⁺TCRV α 7.2⁺CD161^{high} T and innate lymphoid cells in the pregnancies with gestational diabetes, morbidly adherent placenta, and pregnancy hypertension diseases**
Haliloglu Y., Ozcan A., Erdem S., Azizoglu Z. B., Bicer A., Ozarslan Ö. Y., Kilic Ö., Okus F. Z., Demir F., Canatan H., et al.
AMERICAN JOURNAL OF REPRODUCTIVE IMMUNOLOGY, vol.88, no.1, 2022 (SCI-Expanded)
- XXIV. **A novel missense mutation outside the DNAJ domain of DNAJC21 is associated with Shwachman-Diamond syndrome**
Alsavaf M. B., Verboon J. M., DOĞAN M. E., Azizoglu Z. B., Okus F. Z., ÖZCAN A., DÜNDAR M., EKEN A., ALTUNTAŞ H., Sankaran V. G., et al.
BRITISH JOURNAL OF HAEMATOLOGY, vol.197, no.6, 2022 (SCI-Expanded)
- XXV. **Social exclusion and behavior problems in adolescents with cancer and healthy counterparts**
SEZER EFE Y., ÖZBEY H., CANER N., ERDEM E., Kuzucu E. G., KARAKÜKCÜ M., PATIROĞLU T., ÜNAL E., YILMAZ E., ÖZCAN A.
JOURNAL OF PEDIATRIC NURSING-NURSING CARE OF CHILDREN & FAMILIES, vol.64, 2022 (SCI-Expanded)
- XXVI. **The Mutation of CD27 Deficiency Presented With Familial Hodgkin Lymphoma and a Review of the Literature**
Kose D., GÜZELÇİÇEK A., Oz O., Erdem A. Y., Haliloglu Y., Witzel M., Klein C., ÜNAL E.
JOURNAL OF PEDIATRIC HEMATOLOGY ONCOLOGY, vol.44, no.4, 2022 (SCI-Expanded)
- XXVII. **Molecular spectrum of Von Willebrand Disease Type 3**
IŞIK E., AYYILDIZ EMECEN D., Kavaklı K., ŞAHİN F., Durmusalioglu E. A., ALBAYRAK C., SEZGİN EVİM M., ÜNAL E., Belen F. B., Karakurt N., et al.
EUROPEAN JOURNAL OF HUMAN GENETICS, vol.30, no.SUPPL 1, pp.227-228, 2022 (SCI-Expanded)
- XXVIII. **A teenager boy with a novel variant of Sitosterolemia presented with pancytopenia.**
Gok V., Tada H., Ensar Dogan M., ALAKUŞ SARI Ü., ASLAN K., ÖZCAN A., YILMAZ E., KARDAŞ F., KARAKÜKCÜ M., CANATAN H., et al.
Clinica chimica acta; international journal of clinical chemistry, vol.529, pp.61-66, 2022 (SCI-Expanded)
- XXIX. **Evolution and long-term outcomes of combined immunodeficiency due to CARMIL2 deficiency**
Kocamış B., Baser D., Akcam B., Danielson J., Eltan S. B., Haliloglu Y., Sefer A. P., Babayeva R., Akgun G., Charbonnier L., et al.
ALLERGY, vol.77, no.3, pp.1004-1019, 2022 (SCI-Expanded)
- XXX. **Therapeutic effects of vitamin D and IL-22 on methotrexate-induced mucositis in mice**
YILMAZ E., Azizoglu Z. B., Aslan K., Erdem S., Haliloglu Y., Suna P. A., Yay A. H., DENİZ K., TAŞDEMİR A., PER S., et al.
ANTI-CANCER DRUGS, vol.33, no.1, pp.11-18, 2022 (SCI-Expanded)
- XXXI. **A Novel Intronic Mutation Reduces HAX1 Level and is Associated With Severe Congenital**

Neutropenia.

Goktas S., Azizoglu Z. B., Petersheim D., Erdogan M., Eke Gungor H., Bisgin A., Tuğ Bozdoğan S., Eken A., Unal E., Klein C., et al.

Journal of pediatric hematology/oncology, vol.44, 2022 (SCI-Expanded)

XXXII. A Novel Intronic Mutation Reduces HAX1 Level and is Associated With Severe Congenital Neutropenia

GÖKTAŞ S., Azizoglu Z. B., Petersheim D., ERDOĞAN M., Eke Gungor H., Bisgin A., Tug Bozdogan S., EKEN A., ÜNAL E., Klein C., et al.

JOURNAL OF PEDIATRIC HEMATOLOGY ONCOLOGY, vol.44, no.1, 2022 (SCI-Expanded)

XXXIII. Genetic Analysis of a Cohort of 275 Patients with Hyper-IgE Syndromes and/or Chronic Mucocutaneous Candidiasis

Frede N., Rojas-Restrepo J., Caballero Garcia de Oteyza A., Buchta M., Huebscher K., Gamez-Diaz L., Proietti M., Saghafi S., Chavoshzadeh Z., Soler-Palacin P., et al.

JOURNAL OF CLINICAL IMMUNOLOGY, vol.41, no.8, pp.1804-1838, 2021 (SCI-Expanded)

XXXIV. Immunomagnetic separation of B type acute lymphoblastic leukemia cells from bone marrow with flow cytometry validation and microfluidic chip measurements

İçöz K., Eken A., Cinar S., Murat A., Özcan S., Ünal E., Deniz G.

SEPARATION SCIENCE AND TECHNOLOGY, vol.56, no.15, pp.2659-2666, 2021 (SCI-Expanded)

XXXV. Blood repellent superhydrophobic surfaces constructed from nanoparticle-free and biocompatible materials

ÇELİK N., Sahin F., Ruzi M., Yay M., ÜNAL E., ÖNSE S. S.

COLLOIDS AND SURFACES B-BIOINTERFACES, vol.205, 2021 (SCI-Expanded)

XXXVI. Characterization of Th17 and Treg cells in leucocyte adhesion deficiency 1 patients

Erdem S., Haliloglu Y., Haskaloglu S., Arik E., KESKİN Ö., Karadag S. I. K., Eltan S. B., YÜCEL E., AVCILAR H., YILMAZ E., et al.

EUROPEAN JOURNAL OF IMMUNOLOGY, vol.51, pp.298, 2021 (SCI-Expanded)

XXXVII. A nonsense mutation in DIAPH1 gene presents with major T cell defects

Azizoglu Z. B., Okus F. Z., ÖZCAN A., Sawaf B., KÖSE M., CANÖZ Ö., GÜMÜŞ H., Ceylaner S., PATIROĞLU T., ÜNAL E., et al.

EUROPEAN JOURNAL OF IMMUNOLOGY, vol.51, pp.296, 2021 (SCI-Expanded)

XXXVIII. The number and activity of CD3(+)TCR V alpha 7.2(+)CD161(+) cells are increased in children with acute rheumatic fever

Ozkaya M., Baykan A., Cakir M., Vural C., Sunkak S., Unal E., Eken A.

INTERNATIONAL JOURNAL OF CARDIOLOGY, vol.333, pp.174-183, 2021 (SCI-Expanded)

XXXIX. Clinical phenotype, fibrinogen supplementation, and health-related quality of life in patients with afibrinogenemia

Casini A., von Mackensen S., Santoro C., Khayat C. D., Belhani M., Ross C., Dorgalaleh A., Naz A., ÜNAL E., Abdelwahab M., et al.

BLOOD, vol.137, no.22, pp.3127-3136, 2021 (SCI-Expanded)

XL. Type 1 Plasminogen Deficiency With Pulmonary Involvement: Novel Treatment and Novel Mutation

HANGÜL M., TÜZÜNER A. B., Somekh I., Klein C., PATIROĞLU T., ÜNAL E., KÖSE M.

JOURNAL OF PEDIATRIC HEMATOLOGY ONCOLOGY, vol.43, no.4, 2021 (SCI-Expanded)

XLI. Common Variable Immunodeficiency, Autoimmune Hemolytic Anemia, and Pancytopenia Associated With a Defect in IKAROS

Yilmaz E., Kuehn H., Odakir E., Niemela J., Ozcan A., Eken A., Rohlf s M., Cansever M., Gok V., Aydin F., et al.

JOURNAL OF PEDIATRIC HEMATOLOGY ONCOLOGY, no.3, 2021 (SCI-Expanded)

XLII. Refractory and Fatal Presentation of Severe Autoimmune Hemolytic Anemia in a Child With the DNASE1L3 Mutation Complicated With an Additional DOCK8 Variant.

Paç K., Witzel M., Unal E., Rohlf s M., Akyildiz B., Dogan M. E., Poyrazoglu H., Klein C., Patiroglu T.

Journal of pediatric hematology/oncology, vol.43, no.3, 2021 (SCI-Expanded)

XLIII. Comprehensive Mutation Analysis of the RAS/RAF/MEK/ERK Pathway in Paediatric Leukaemia and

Significant Inferences

Akin-Bali D. F., AKTAŞ S. H., ÖZCAN A., YILMAZ E., AYDIN F., GÖK V., ÜNAL E., KARAKÜKCÜ M.
HONG KONG JOURNAL OF PAEDIATRICS, vol.26, no.2, pp.75-87, 2021 (SCI-Expanded)

- XLIV. **Effect of vitamin K2 and vitamin D3 on bone mineral density in children with acute lymphoblastic leukemia: a prospective cohort study.**
SOLMAZ I., ÖZDEMİR M. A., ÜNAL E., Abdurrezzak Ü., Muhtaroglu S., KARAKÜKCÜ M.
Journal of pediatric endocrinology & metabolism : JPEM, vol.34, no.4, pp.441-447, 2021 (SCI-Expanded)
- XLV. **The Spectrum of Underlying Diseases in Children with Torticollis**
Tumturk A., Gorkem S. B., Ozmansur E. N., Peduk Y., Arslan U., Gok V., Dogan H., Cetin B. Ş., Sahin A., Gumus H., et al.
TURKISH NEUROSURGERY, no.3, pp.389-398, 2021 (SCI-Expanded)
- XLVI. **Expanded circulating hematopoietic stem/progenitor cells as novel cell source or the treatment of TCIRG1 osteopetrosis**
Capo V., Penna S., Merelli I., Barcella M., Scala S., Basso-Ricci L., Draghici E., Palagano E., Zonari E., Desantis G., et al.
HAEMATOLOGICA, vol.106, no.1, pp.74-86, 2021 (SCI-Expanded)
- XLVII. **Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency**
ÖZEN A. O., ÖZEN A. O., Edwards E. S. J., Pillay B., ÖZEN A. O., ÖZEN A. O., ÖZEN A. O., Zoghi S., Abolhassani H., et al.
BLOOD, vol.136, no.23, pp.2638-2655, 2020 (SCI-Expanded)
- XLVIII. **Hepatitis-associated aplastic anemia in pediatric patients: single center experience.**
Altay D., Yilmaz E., Özcan A., Karakükçü M., Ünal E., Arslan D.
Transfusion and apheresis science : official journal of the World Apheresis Association : official journal of the European Society for Haemapheresis, vol.59, 2020 (SCI-Expanded)
- XLIX. **Different Clinical Presentation of 3 Children With Familial Hemophagocytic Lymphohistiocytosis With 2 Novel Mutations.**
Akyol S., Ozcan A., Sekine T., Chiang S., Yilmaz E., Karakurkcü M., Patiroglu T., Bryceson Y., Unal E.
Journal of pediatric hematology/oncology, vol.42, 2020 (SCI-Expanded)
- L. **Microfluidic Chip based direct triple antibody immunoassay for monitoring patient comparative response to leukemia treatment**
İÇÖZ K., Akar U., ÜNAL E.
BIOMEDICAL MICRODEVICES, vol.22, no.3, 2020 (SCI-Expanded)
- LI. **Whole exome sequencing (WES) approach for diagnosing primary immunodeficiencies (PIDs) in a highly consanguineous community**
Simon A. J., Golan A. C., Lev A., Stauber T., Barel O., Somekh I., Klein C., AbuZaitun O., Eyal E., Kol N., et al.
CLINICAL IMMUNOLOGY, vol.214, 2020 (SCI-Expanded)
- LII. **Role of a second transplantation for children with acute leukemia following posttransplantation relapse: a study by the Turkish Bone Marrow Transplantation Study Group**
Hazar V., Karasu G. T., Uygun V., Ozbek N., KARAKÜKCÜ M., Ozturk G., Daloglu H., Kilic S. C., Aksu T., ÜNAL E., et al.
LEUKEMIA & LYMPHOMA, vol.61, no.6, pp.1465-1474, 2020 (SCI-Expanded)
- LIII. **ILC3 deficiency and generalized ILC abnormalities in DOCK8-deficient patients**
Eken A., Cansever M., Okus F. Z., Erdem S., Nain E., Azizoglu Z. B., Haliloglu Y., Karakukcu M., Ozcan A., Devecioglu O., et al.
ALLERGY, vol.75, no.4, pp.921-933, 2020 (SCI-Expanded)
- LIV. **Congenital afibrinogenemia in a 4-year-old girl complicated with acute lymphoblastic leukemia**
ÖZCAN A., Samur B., AKYOL Ş., Erdogmus N. A., PATIROĞLU T., KARAKÜKCÜ M., ÜNAL E.
TURKISH JOURNAL OF PEDIATRICS, vol.62, no.2, pp.289-292, 2020 (SCI-Expanded)
- LV. **Twenty children with non-Wilms renal tumors from a reference center in Central Anatolia, Turkey**
Ünal E., Yilmaz E., Özcan A., Işık B., Karakükçü M., Turan C., Akgün H., Öztürk F., Coşkun A., Özdemir M., et al.
Turkish journal of medical sciences, vol.50, pp.18-24, 2020 (SCI-Expanded)
- LVI. **Human FCH01 deficiency reveals role for clathrin-mediated endocytosis in development and function of T cells**

- Lyszkiwicz M., Zietara N., Frey L., Pannicke U., Stern M., Liu Y., Fan Y., Puchalka J., Hollizeck S., Somekh I., et al. NATURE COMMUNICATIONS, vol.11, no.1, 2020 (SCI-Expanded)
- LVII. Factor 8 Gene Mutation Spectrum of 270 Patients with Hemophilia A: Identification of 36 Novel Mutations**
ATİK T., IŞIK E., Onay H., Akgun B., Shamsali M., Kavaklo K., SEZGİN EVİM M., TÜYSÜZ KINTRUP G., Ozbek N. Y., ŞAHİN F., et al.
TURKISH JOURNAL OF HEMATOLOGY, vol.37, no.3, pp.145-153, 2020 (SCI-Expanded)
- LVIII. Assesment of Patients with Von Willebrand Disease with ISTH/BAT and PBQ Scores**
Belen A., Ümit E., Zengin Y., Sezgin E., Ünal E., Özbaş H., Acipayam C.
Turkish journal of haematology : official journal of Turkish Society of Haematology, vol.37, pp.57-58, 2020 (SCI-Expanded)
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